

Table 4: Gag

MAb ID	HXB2 Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)
122 32/5.8.42	Gag(dis) References: [Papsidero (1989)] • 32/5.8.42: Inhibited infectivity of cell free virus – bound to both peptides, ELDRWEKI and ALDKIE –Papsidero89	p17(12-19 + 100–105 IIIB)	ELDRWEKI + ALDKIE	no	Viral lysate	murine(IgG)
123 32/5.8.42	Gag(dis) References: [Papsidero (1989)] • 32/5.8.42: Inhibited infectivity of cell free virus – bound ELDRWEKI and ALDKIE –Papsidero89	p17(12-19 + 100–105 IIIB)	ELDRWEKI + ALDKIE	no	Viral lysate	murine(IgG)
124 CH9B2	Gag() Donor: R. B. Ferns and R. S. Tedder References: [Ferns (1987), Ferns (1989)] • CH9B2: Reactive against p18 and p55 –Ferns87 • CH9B2: UK Medical Research Council AIDS reagent: ARP349	p17()			Inact CBL-1	murine(IgG ₁)
125 G11G1	Gag() References: [Shang (1991), Pincus (1996)] • G11G1: Immunotoxins were generated by linking Env MAbs to ricin A – immunotoxins mediated cell killing, but only if the antigen was expressed at the cell surface – ricin-G11G1 did not mediate cell killing –Pincus96	p17()			?	rat()
126 2A6	Gag() Donor: A. O. Arthur, Frederick Cancer Research and Development Center, Frederick, MD References: [Pincus (1998)] • 2A6: Part of a panel of 17 MAbs used as controls testing for the dual specificity of MAb G11H3 for both p17 and mycoplasma –Pincus98	p17()			?	()
127 G11H3	Gag(dis) References: [Shang (1991), Pincus (1998)] • G11H3: This MAb is cross-reactive between p17 and mycoplasma – this antibody binds strain specifically to the variable lipoprotein (Vlp) F of <i>M. hyorhinis</i> , in the region of the carboxy-terminal repeat CGGSTPTPEQGNNQG-GSTPTPEQGNSQVSK – the p17 epitope is discontinuous, but p17 and VlpF share the tetrapeptide SQVS –Pincus98	p17(Gag dis)			?	()

HIV Monoclonal Antibodies

MAb ID	HXB2 Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)
128 HyHIV-19	Gag(dis) References: [Liu (1995), Ota (1998)]	p17(dis JMH1)		no	rec p17	murine(IgG ₁)
	<ul style="list-style-type: none"> HyHIV-19: Does not react with p17 peptides – Ka is $3.7 \times 10^6 \text{ M}^{-1}$ for rec p17 – inhibited growth of HIV-1 JMH1 in MT-4 cells when added 24 hours after the initial culture –Ota98a 					
129 LH-104-A	Gag(dis) References: [Haaheim (1991)]	p24(dis BRU)	DIRQGP + QGVGGP	no	104 amino acid peptide	murine(IgG ₁ κ)
	<ul style="list-style-type: none"> LF-104-A: Hexapeptide scans revealed two reactive p24 peptides – cross-competition studies indicated the region 270–286 –Haaheim91 LH-104-A: UK Medical Research Council AIDS reagent: ARP307 					
130 EH12E1	Gag(dis) Donor: R. B. Ferns and R. S. Tedder References: [Ferns (1987), Ferns (1989)]	p24(Gag dis)			Inact CBL-1	murine(IgG ₁)
	<ul style="list-style-type: none"> EH12E1: Reacted with p55 and p24 in WB –Ferns87 EH12E1: UK Medical Research Council AIDS reagent: ARP313 					
131 LH-104-C	Gag(dis) References: [Haaheim (1991)]	p24(dis BRU)	GPKEPF + QGVGGP	no	104 amino acid peptide	murine(IgG ₃ κ)
	<ul style="list-style-type: none"> LF-104-C: Hexapeptide scans revealed two reactive p24 peptides – cross-competition studies indicated the region 351–373 –Haaheim91 LH-104-C: UK Medical Research Council AIDS reagent: ARP309 					
132 71-31	Gag() References: [Gorny (1989), Robinson (1990b), Robinson (1991), Spear (1993), Gorny (1997), Gorny (1998), Bandres (1998)]	p24()		no	HIV-1	human(IgG ₁ λ)
	<ul style="list-style-type: none"> 71-31: Did not enhance HIV-1 IIIB infection –Robinson90a 71-31: No enhancing or neutralizing activity –Robinson91 71-31: Did not mediate deposition of complement component C3 on HIV infected cells –Spear93 71-31: Included as a negative control in studies that demonstrate that CXCR4 can bind to gp120 in the absence of CD4-gp120 interactions, and that this binding can be enhanced by Env deglycosylation –Bandres98 71-31: NIH AIDS Research and Reference Reagent Program: 530 					

HIV Monoclonal Antibodies

MAb ID	HXB2 Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)
133 V7-8	Gag()	p24()		no	HIV-1 infection	murine(IgG ₃ κ)
References: [Robinson (1990b), Montefiori (1991)] <ul style="list-style-type: none"> • V7-8: Did not enhance HIV-1 IIIB infection –Robinson90a • V7-8: Reacted with HIV-1IIIB, RF, and MN –Montefiori91 • V7-8: NIH AIDS Research and Reference Reagent Program: 381 						
134 98-4.9	Gag()	p24()		no	HIV-1 infection	murine(IgG ₃ λ)
References: [Gorny (1989)]						
135 98-4.3	Gag()	p24()		no	HIV-1 infection	human(IgG ₁ λ)
References: [Robinson (1991)] <ul style="list-style-type: none"> • 98-4.3: No enhancing or neutralizing activity –Robinson91 						
136 IE8G2	Gag()	p24()			Inact CBL-1	murine(IgG ₁)
Donor: R. B. Ferns and R. S. Tedder References: [Ferns (1987), Ferns (1989)] <ul style="list-style-type: none"> • IE8G2: Reacted with both p55 and p24 – broadly reactive – showed less than 75% homologous inhibition –Ferns87 • IE8G2: UK Medical Research Council AIDS reagent: ARP347 						
137 human sera	Gag()	p24()			HIV-1 infection	human(IgG)
References: [Binley (1997b)] <ul style="list-style-type: none"> • Retention of anti-Env antibodies and loss of anti-Gag antibodies during progression was studied, and suggested to be the result of the loss of T-cell help and the unique ability of Env to stimulate B cells even in a backdrop of declining CD4 cells, because of the ability of Env to bind to the CD4 molecule –Binley97a 						
138 241-D	Gag()	p24()		no		human(IgG ₁ λ)
Donor: Susan Zolla-Pazner (NYU Med. Center) References: [Gorny (1989), Tyler (1990), Robinson (1991)] <ul style="list-style-type: none"> • 241-D: An antibody by this name is available in the NIH AIDS Research and Reference Reagent Program, and they refer to the papers: Gorny89,Tyler90,Robinson91, but no p24 MAb by this name is discussed in these papers • 241-D: MH AIDS Research and Reference Reagent program: 1244 						

B Cell

HIV Monoclonal Antibodies

MAb ID	HXB2 Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)
139 183-H12-5C	Gag()	p24()		no	unk	murine(IgG ₁)
<p>Donor: Bruce Chesebro and Kathy Wehrly, Rocky Mountain Laboratories, Hamilton, Montana</p> <p>References: [Chesebro (1992), Toohey (1995), Wehrly & Chesebro(1997)]</p> <ul style="list-style-type: none"> • 183-H12-5C: Cross-reacts with HIV1 and HIV-2 p24, and SIV p27 • 183-H12-5C: Used as antigen capture reagent for p24 ELISA –Chesebro92,Toohey95 • 183-H12-5C: Cross-reacts with HIV1 and HIV-2 p24, and SIV p27 –Wehrly97 • 183-H12-5C: NIH AIDS Research and Reference Reagent Program: 3537 						
140 ED8	Gag(dis)	p7(Gag dis)		no	purified NCp7	murine(IgG)
<p>References: [Tanchou (1995)]</p> <ul style="list-style-type: none"> • ED8: Binds NCp7 independent of Zn fingers, does not react with NCp15 –Tanchou95 						
141 AC2	Gag(dis)	p7(Gag dis)		no	purified NCp7	murine(IgG)
<p>References: [Tanchou (1995)]</p> <ul style="list-style-type: none"> • AC2: Binds NCp7 independent of Zn fingers, does not react with NCp15 –Tanchou95 						
142 CD9	Gag(dis)	p7(Gag dis)		no	purified NCp7	murine(IgG)
<p>References: [Tanchou (1995)]</p> <ul style="list-style-type: none"> • CD9: Binds NCp7 independent of Zn fingers, does not react with NCp15 –Tanchou95 						
143 BE10	Gag(dis)	p7(Gag dis)		no	purified NCp7	murine(IgG)
<p>References: [Tanchou (1995)]</p> <ul style="list-style-type: none"> • BE10: Binding NCp7 requires Zn fingers, does not react with NCp15, inhibits NCp7-tRNA interaction –Tanchou95 						